

OSBORNE et al.
Appl. No. 10/567,453
Atty. Ref.: 620-412
Amendment
February 7, 2011

AMENDMENTS TO THE CLAIMS:

Please amend the claims as follows:

1. (Currently Amended) A method for the in vitro culture of a myeloma cell line which comprises:

(a) inoculating a culture medium with the myeloma cell line, said medium being capable of supporting the growth of said myeloma cell line and comprising iron at concentrations in the medium of from about 0.064 mg/L to about 1.6 mg/L, wherein said medium does not contain transferrin, a lipophilic chelator, a synthetic nitrogen-containing chelator or a lipophilic synthetic nitrogen-containing chelator ; and

(b) growth of the inoculated culture medium under appropriate conditions and using agitated suspension culture,and

wherein the source of iron is a soluble iron compound selected from the group consisting of ferric ammonium citrate, ferric ammonium oxalate, ferric ammonium fumarate, ferric ammonium malate and ferric ammonium succinate.

Claim 2. (Cancelled)

Claim 3. (Cancelled)

4. (Original) The method of claim 1 wherein the concentration of iron in the medium is from about 0.16 mg/L to about 0.32 mg/L.

Claims 5-8. (Canceled)

9. (Currently Amended) The method of claim [[7]]1 wherein the source of iron ferric ammonium compound is ferric ammonium citrate.

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10. (Currently Amended) A method for the in vitro culture of a myeloma cell line which comprises:

(a) inoculating a culture medium with the myeloma cell line, said medium being capable of supporting the growth of said myeloma cell line and comprising ferric ammonium citrate at a concentration in the medium of from about 0.4 mg/L to about 10 mg/L, wherein said medium does not contain transferrin, a lipophilic chelator, a synthetic nitrogen-containing chelator or a lipophilic synthetic nitrogen- containing chelator; and

(b) growth of the inoculated culture medium under appropriate conditions and using agitated suspension culture.

Claim 11. (Cancelled)

Claim 12. (Cancelled)

13. (Original) The method of claim 10 wherein the ferric ammonium citrate is present in the medium at a concentration of from about 1 mg/L to about 2 mg/L.

14. (Previously Presented) The method of claim 1 wherein the medium is serum free, protein free, free of components of animal derivation or is chemically defined.

15. (Previously Presented) The method of claim 1 wherein the myeloma cell line is selected from the group consisting of an NSO series cell line, a P3 series cell line, MOPC series cell line, the MPC-11 cell line, the J558L cell line, the K6H6/B5 cell line, the 45.6.TG1.7 cell line, the YO cell line, the Y3 HTK cell line, the RPMI 8226 cell line and the U266B1 cell line.

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16. (Previously Presented) The method of claim 1 wherein the myeloma cell line is the NSO cell line.

Claims 17-32. (Cancelled)

33. (Currently Amended) A process for obtaining a mammalian cell product comprising culturing a myeloma cell capable of producing said product under agitated suspension culture and in a culture medium capable of supporting the growth of said myeloma cell line, said medium comprising iron at concentrations in the medium of from about 0.064 mg/L to about 1.6 mg/L, or ferric ammonium citrate at a concentration in the medium of from about 0.4 mg/L to about 10 mg/L, wherein said medium does not contain transferrin, a lipophilic chelator, a synthetic nitrogen- containing chelator or a lipophilic synthetic nitrogen-containing chelator; and recovering said mammalian cell product, and

wherein the source of iron is a soluble iron compound selected from the group consisting of ferric ammonium citrate, ferric ammonium oxalate, ferric ammonium fumarate, ferric ammonium malate and ferric ammonium succinate.

Claim 34. (Cancelled)

Claim 35. (Cancelled)

36. (Original) The process of claim 33 wherein the concentration of iron in the medium is from about 0.16 mg/L to about 0.32 mg/L.

Claims 37-40. (Cancelled)

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41. (Currently Amended) The process of claim 40 wherein the source of iron
~~ferric ammonium compound~~ is ferric ammonium citrate.

Claim 42. (Cancelled)

Claim 43. (Cancelled)

Claim 44. (Cancelled)

45. (Previously Presented) The process of claim 33 wherein the ferric ammonium citrate is present in the medium at a concentration of from about 1 mg /L to about 2 mg /L.

46. (Previously Presented) The process of claim 33 wherein the medium is serum free, protein free, free of components of animal derivation or is chemically defined.

47. (Previously Presented) The process of claim 33 wherein the myeloma cell line is selected from the group consisting of an NSO series cell line, a P3 series cell line, a MOPC series cell line, the MPC-11 cell line, the J558L cell line, the K6H6/B5 cell line, the 45.6.TG1.7 cell line, the YO cell line, the Y3 HTK cell line, the RPMI 8226 cell line and the U266B1 cell line.

48. (Previously Presented) The process of claim 33 wherein the myeloma cell line is the NSO cell line.

49. (Previously Presented) The process of claim 33 wherein the cell product is selected from the group consisting of polypeptides, proteins, hormones, lymphokines, interleukins and industrially and therapeutically useful enzymes.

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50. (Original) The process of claim 49 wherein the cell product is an antibody or fragment thereof.